

Voluntary Industry Reporting Form for 6(a)(2) Adverse Effects Incident Information

-001

Provide all known, required information. If required data field information is unknown, designate as such in appropriate area. Page # 1 of 3

Row 1	Reporter name: [REDACTED]	Submission date:	Contact person (if different than reporter)	Internal ID 1-49655563
Administrative Data	Address: Ontario		Address:	
	Phone #: [REDACTED]		Phone #:	
	Incident Status: New	Location and date of incident Ontario 08/27/2017	Date registrant became aware of incident: 9/10/2017	Was incident part of larger study?
Row 2	EPA Registration # (Product 1) 24359	EPA Registration # (Product 2)	EPA Registration # (Product 3)	
Pesticide(s) Involved	A.I. (s)	A.I. (s)	A.I. (s)	
	Product 1 Name Glyfos Soluble Concentrate Herbicide Canada	Product 2 Name	Product 3 Name	
	Exposed to concentrate prior to dilution? NA	Exposed to concentrate prior to dilution?	Exposed to concentrate prior to dilution?	
	Formulation	Formulation	Formulation	
Row 3	Evidence label directions were not followed? No Intentional misuse? No	Incident site: (examples include home, yard, school, industrial, nursery/greenhouse, surface water, commercial turf, building/office, forest/ woods, agricultural (specify crop) right-of-way (rail, utility, highway)) Own Residence	Situation: (act of using product): (examples include mixing/loading, reentry, application, transportation, repair/ maintenance of application equipment, manufacturing/ formulating) See Description Notes	
Incident Circumstances	Applicator certified PCO? Not applicable			
	How exposed: (examples include direct contact with treated surface, ingestion, spill, drift, runoff) See Incident Description			

Personal privacy information

*9/10/2017 10:05:58 AM Glyphos Soluble Concentrate Herbicide Canada
PCP #24359*

HX: The caller got some of the product on his hand 2 weeks ago. He did take a shower after exposure. Shortly after exposure he started to experience joint and muscle pain. Could his symptoms be related to this product?

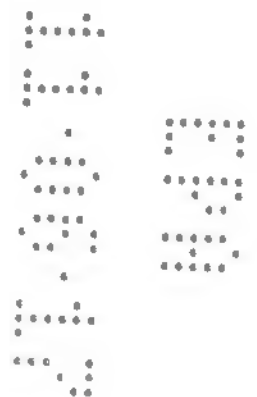
A: The symptoms described would not be expected from exposure to this product.

- Product may cause temporary skin irritation.*
- Consider other causes for your symptoms.*
- Please call back with any further questions or concerns.*

Consulted with AB

10/6/2017 11:48:10 AM PMRA report generated.

10/11/2017 9:44:57 PM PMRA report sent.



Voluntary Industry Reporting Form for 6(a)(2) Incident Information Involving Humans

Provide all known, required information. If required data field information is unknown, designate as such in appropriate area. Page # 3 of 3

Demographic information Age: <i>Unknown</i> Sex: <i>Male</i> Occupation: (if relevant)	Exposure route: <i>Dermal</i>	Was adverse effect result of suicide/homicide or attempted suicide/homicide? <i>No</i>	Was protective clothing worn (specify)? <i>Not applicable</i>
If female, pregnant? <i>Did not query</i>	Was exposure occupational? <i>No</i> If yes, days lost due to illness:	Time between exposure and onset of symptoms: <i>See Symptoms</i>	
Type of medical care sought: (examples include none, clinic, hospital emergency department, private physician, PCC, hospital inpatient). <i>On-site</i>	List signs/symptoms/adverse effects. <i>Joint pain, 24 hrs or less;</i> <i>Other miscellaneous - Muscle pain, 24 hrs or less;</i>		If lab tests were performed, list test names and results (If available, submit reports). <i>Not Reported</i>
Exposure data: Amount of pesticide: Exposure duration: Weight:			
Human severity category: <i>HC</i>			

This box can be used to provide any explanatory or qualifying information surrounding the incident. (add additional pages if necessary)

Internal ID #
1-49655563